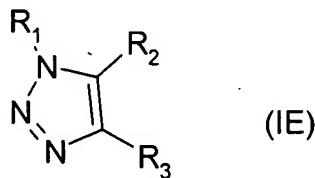


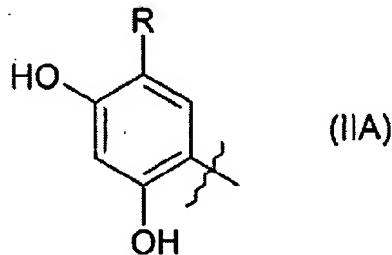
Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Currently Amended) A compound of formula (IE) or a salt, or N-oxide, or hydrate or solvate thereof, for use in human or veterinary medicine:



wherein R₁ has the formula (IIA):



wherein R represents bromo, chloro, phenyl, C₁-C₆ alkyl or phenyl(C₁-C₆ alkyl)

R₂ is hydrogen or

a carboxamide radical; or

a non aromatic carbocyclic or non aromatic heterocyclic ring wherein a ring carbon is optionally substituted, and/or a ring nitrogen is optionally substituted by a group of formula -(Alk¹)_p-(Z)_r(Alk²)_s-Q wherein in any compatible combination Alk¹ and Alk² are divalent C₁-C₆ alkylene or C₂-C₆ alkenylene radicals,

p, r and s are independently 0 or 1,

Z is -O-, -S-, -(C=O)-, -(C=S)-, -SO₂-, -C(=O)O-, -C(=O)NR^A-, -C(=S)NR^A-, -SO₂NR^A-, -NR^AC(=O)-, -NR^ASO₂- or -NR^A- wherein R^A is hydrogen or C₁-C₆ alkyl, and

Q is hydrogen or a carbocyclic or heterocyclic radical;

wherein each of Alk¹, Alk², and Q are optionally substituted with one or more substituents selected from (C₁-C₆)alkyl, (C₁-C₆)alkoxy, hydroxy, hydroxy(C₁-C₆)alkyl, mercapto, mercapto(C₁-C₆)alkyl, (C₁-C₆)alkylthio, halo (including fluoro and chloro), trifluoromethyl, trifluoromethoxy, nitro, nitrile (-CN), oxo, phenyl, -COOH, -COOR^A, -COR^A, -SO₂R^A, -CONH₂, -SO₂NH₂, -CONHR^A, -SO₂NHR^A, -CONR^AR^B, -SO₂NR^AR^B, -NH₂, -NHR^A, -NR^AR^B, -OCONH₂, -OCONHR^A, -OCONR^AR^B, -NHCOR^A, -NHCOOR^A, -NR^BCOOR^A, -NHSO₂OR^A, -NR^BSO₂OR^A, -NHCONH₂, -NR^ACONH₂, -NHCONHR^B, -NR^ACONHR^B, -NHCONR^AR^B, or -NR^ACONR^AR^B, wherein R^A and R^B are independently a (C₁-C₆)alkyl group; and

R₃ is hydrogen, optionally substituted cycloalkyl, cycloalkenyl, C₁-C₆ alkyl, C₂-C₆ alkenyl, or C₂-C₆ alkynyl; or a carboxyl, carboxamide or carboxyl ester group,

PROVIDED THAT at least one of R₂ and R₃ is present and is other than hydrogen.

Claims 2-14 (Cancelled)

15. (Currently Amended) The compound as claimed in claim 1 wherein R₂ is phenyl, 2-, 3-, or 4-pyridyl, 2- or 3-furanyl, 2- or 3-thienyl, or thiazolyl, optionally substituted by one or more of methoxy, ethoxy, methylenedioxy, ethylenedioxy, fluoro, chloro, bromo, or trifluoromethyl.

16. (Cancelled)

17. (Withdrawn) The compound as claimed in claim 1 wherein R₂ is a carboxamide radical of formula -CONR^B(Alk)_nR^A wherein

Alk is an optionally substituted divalent alkylene, alkenylene or alkynylene radical,

n is 0 or 1,

R^B is hydrogen or a C₁-C₆ alkyl or C₂-C₆ alkenyl group,

R^A is hydroxy or an optionally substituted carbocyclic or heterocyclic ring,

or R^A and R^B taken together with the nitrogen to which they are attached form an N-heterocyclic ring which may optionally contain one or more additional hetero atoms selected from O, S and N, and which may optionally be substituted on one or more ring C or N atoms.

18. (Withdrawn) The compound as claimed in claim 17 wherein

Alk is an optionally substituted -CH₂-, -CH₂CH₂-, -CH₂CH₂CH₂-, CH₂CH=CH-, or -CH₂CCCH₂- radical.

n is 0 or 1 ,

R^B is hydrogen, methyl, ethyl, n- or iso-propyl, or allyl,

R^A is hydroxy, hydroxy and/or chloro-substituted phenyl, 3,4 methylenedioxyphenyl, pyridyl, furyl, thienyl, N-piperazinyl, or Nmorpholinyl,

or R^A and R^B taken together with the nitrogen to which they are attached form a morpholino, piperidinyl, piperazinyl or N-phenylpiperazinyl ring.

19. (Withdrawn) The compound as claimed in claim 17 wherein n is 0, R^B is hydrogen and R^A is hydroxy or an optionally substituted carbocyclic or heterocyclic ring.

20. (Withdrawn) The compound as claimed in claim 1 wherein R₃ is hydrogen, methyl, ethyl, n- or iso-propyl, trifluoromethyl, or hydroxyethyl.

21. (Withdrawn) The compound as claimed in claim 1 wherein R₃ is a carboxamide group -CONR^B(Alk)_nR^A wherein
Alk is an optionally substituted divalent alkylene, alkenylene or alkynylene radical,
n is 0 or 1,
R^B is hydrogen or a C₁-C₆ alkyl or C₂-C₆ alkenyl group,
R^A is hydroxy or an optionally substituted carbocyclic or heterocyclic ring,
or R^A and R^B taken together with the nitrogen to which they are attached form an N-heterocyclic ring which may optionally contain one or more additional hetero atoms selected from O, S and N, and which may optionally be substituted on one or more ring C or N atoms.

22. (Withdrawn – Currently Amended) A method of treatment of diseases or conditions mediated by excessive or inappropriate HSP90 activity in mammals which method comprises administering to the mammal an amount of a compound of formula (IE) as defined in claim 1-, or a salt, ~~hydrate or solvate~~ thereof, effective to inhibit said HSP90 activity.

23. (Withdrawn – Currently Amended) The method as claimed in claim 22 for treatment of immunosuppression; or the treatment of cancer; viral disease; diseases; inflammatory diseases such as rheumatoid arthritis, asthma, multiple sclerosis, Type I diabetes, lupus, psoriasis and inflammatory bowel disease; cystic fibrosis; or angiogenesis-related diseases ~~disease such as diabetic retinopathy, haemangiomas, and endometriosis; or treatment~~ for protection of normal cells against chemotherapy-induced toxicity; or

treatment of diseases where failure to undergo apoptosis is an underlying factor; or
treatment for protection from hypoxia-ischemic injury due to elevation of Hsp70 in the
heart and brain; or treatment of scrapie/CJD, Huntingdon's disease or and Alzheimer's
disease.

24. (Cancelled)

25. (Currently Amended) A pharmaceutical or veterinary composition comprising a compound as defined in claim 1, or a salt ~~hydrate or solvate~~ thereof, together with a pharmaceutically or veterinarianily acceptable carrier.